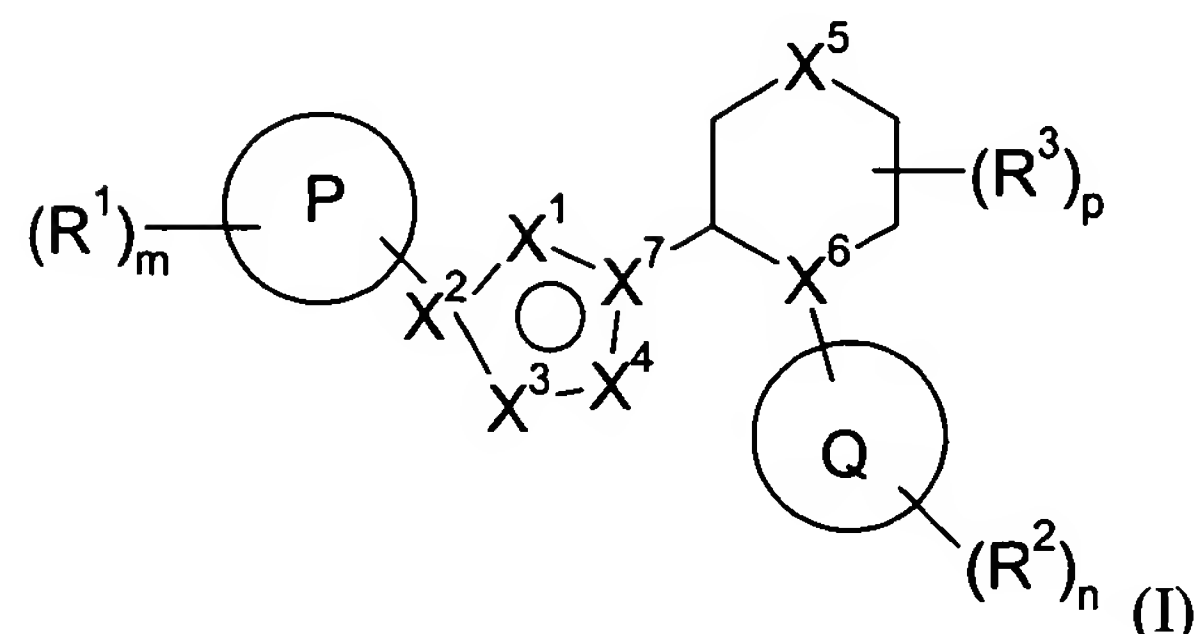


AMENDMENTS TO THE CLAIMS

1. (Original) A compound according to formula I



wherein

P is selected from aryl and heteroaryl;

R^1 is attached to P via a carbon atom on ring P and is selected from the group consisting of hydroxy, halo, nitro, C_{1-6} alkylhalo, OC_{1-6} alkylhalo, C_{1-6} alkyl, OC_{1-6} alkyl, C_{2-6} alkenyl, OC_{2-6} alkenyl, C_{2-6} alkynyl, OC_{2-6} alkynyl, C_{0-6} alkyl C_{3-6} cycloalkyl, OC_{0-6} alkyl C_{3-6} cycloalkyl, C_{0-6} alkylaryl, OC_{0-6} alkylaryl, CHO, $(CO)R^5$, $O(CO)R^5$, $O(CO)OR^5$, $O(CNR^5)OR^5$, C_{1-6} alkyl OR^5 , OC_{2-6} alkyl OR^5 , C_{1-6} alkyl $(CO)R^5$, OC_{1-6} alkyl $(CO)R^5$, C_{0-6} alkyl CO_2R^5 , OC_{1-6} alkyl CO_2R^5 , C_{0-6} alkylcyano, OC_{2-6} alkylcyano, C_{0-6} alkyl NR^5R^6 , OC_{2-6} alkyl NR^5R^6 , C_{1-6} alkyl $(CO)NR^5R^6$, OC_{1-6} alkyl $(CO)NR^5R^6$, C_{0-6} alkyl $NR^5(CO)R^6$, OC_{2-6} alkyl $NR^5(CO)R^6$, C_{0-6} alkyl $NR^5(CO)NR^5R^6$, C_{0-6} alkyl SR^5 , OC_{2-6} alkyl SR^5 , C_{0-6} alkyl $(SO)R^5$, OC_{2-6} alkyl $(SO)R^5$, C_{0-6} alkyl SO_2R^5 , OC_{2-6} alkyl SO_2R^5 , C_{0-6} alkyl $(SO_2)NR^5R^6$, OC_{2-6} alkyl $(SO_2)NR^5R^6$, C_{0-6} alkyl $NR^5(SO_2)R^6$, OC_{2-6} alkyl $NR^5(SO_2)R^6$, C_{0-6} alkyl $NR^5(SO_2)NR^5R^6$, OC_{2-6} alkyl $NR^5(SO_2)NR^5R^6$, $(CO)NR^5R^6$, $O(CO)NR^5R^6$, NR^5OR^6 , C_{0-6} alkyl $NR^5(CO)OR^6$, OC_{2-6} alkyl $NR^5(CO)OR^6$, SO_3R^5 and a 5- or 6-membered ring containing atoms independently selected from the group consisting of C, N, O and S;

X^1 is selected from the group consisting of: N, NR^4 and CR^4 ;

X^2 is selected from the group consisting of: C and N;

X^3 is selected from the group consisting of: CR^4 , N and O;

X^4 is selected from the group consisting of: CR^4 , N, NR^4 and O;

X^5 is selected from the group consisting of: a bond, $CR^4R^{4'}$, NR^4 , O, S, SO and SO_2 ;

X^6 is selected from the group consisting of: CR^4 and N;

X^7 is selected from the group consisting of: C and N;

R^4 is independently selected from a group consisting of hydrogen, hydroxy, C_{1-6} alkyl, C_{0-6} alkylcyano, oxo, $=NR^5$, $=NOR^5$, C_{1-4} alkylhalo, halo, C_{3-7} cycloalkyl, $O(CO)C_{1-4}$ alkyl, C_{1-4} alkyl(SO) C_{0-4} alkyl, C_{1-4} alkyl(SO_2) C_{0-4} alkyl, (SO) C_{0-4} alkyl, (SO_2) C_{0-4} alkyl, OC_{1-4} alkyl, C_{1-4} alkylOR⁵ and C_{0-4} alkylNR⁵R⁶;

Q is selected the group consisting of heterocycloalkyl and heteroaryl;

R^2 and R^3 are independently selected from the group consisting of: hydroxy, C_{0-6} alkylcyano, oxo, $=NR^5$, $=NOR^5$, C_{1-4} alkylhalo, halo, C_{1-6} alkyl, C_{3-6} cycloalkyl, C_{0-6} alkylaryl, C_{0-6} alkylheteroaryl, C_{1-6} alkylcycloalkyl, C_{0-6} alkylheterocycloalkyl, OC_{1-4} alkyl, OC_{0-6} alkylaryl, $O(CO)C_{1-4}$ alkyl, $(CO)OC_{1-4}$ alkyl, C_{0-4} alkyl(S) C_{0-4} alkyl, C_{1-4} alkyl(SO) C_{0-4} alkyl, C_{1-4} alkyl(SO_2) C_{0-4} alkyl, (SO) C_{0-4} alkyl, (SO_2) C_{0-4} alkyl, C_{1-4} alkylOR⁵, C_{0-4} alkylNR⁵R⁶ and a 5- or 6-membered ring containing atoms independently selected from C, N, O and S, which ring may optionally be fused with a 5- or 6-membered ring containing atoms independently selected from the group consisting of C, N and O and wherein said ring and said fused ring may be substituted by one or more A;

wherein any C₁₋₆alkyl, aryl, or heteroaryl defined under R¹, R² and R³ may be substituted by one or more A ;

A is selected from the group consisting of: hydrogen, hydroxy, halo, nitro, oxo, C₀₋₆alkylcyano, C₀₋₄alkylC₃₋₆cycloalkyl, C₁₋₆alkyl, -OC₁₋₆alkyl, C₁₋₆alkylhalo, OC₁₋₆alkylhalo, C₂₋₆alkenyl, C₀₋₃alkylaryl, C₀₋₆alkylOR⁵, OC₂₋₆alkylOR⁵, C₀₋₆alkylSR⁵, OC₂₋₆alkylSR⁵, (CO)R⁵, O(CO)R⁵, OC₂₋₆alkylcyano, OC₁₋₆alkylCO₂R⁵, O(CO)OR⁵, OC₁₋₆alkyl(CO)R⁵, C₁₋₆alkyl(CO)R⁵, NR⁵OR⁶, C₀₋₆NR⁵R⁶, OC₂₋₆alkylNR⁵R⁶, C₀₋₆alkyl(CO)NR⁵R⁶, OC₁₋₆alkyl(CO)NR⁵R⁶, OC₂₋₆alkylNR⁵(CO)R⁶, C₀₋₆alkylNR⁵(CO)R⁶, C₀₋₆alkylNR⁵(CO)NR⁵R⁶, O(CO)NR⁵R⁶, C₀₋₆alkyl(SO₂)NR⁵R⁶, OC₂₋₆alkyl(SO₂)NR⁵R⁶, C₀₋₆alkylNR⁵(SO₂)R⁶, OC₂₋₆alkylNR⁵(SO₂)R⁶, SO₃R⁵, C₁₋₆alkylNR⁵(SO₂)NR⁵R⁶, OC₂₋₆alkyl(SO₂)R⁵, C₀₋₆alkyl(SO₂)R⁵, C₀₋₆alkyl(SO)R⁵, OC₂₋₆alkyl(SO)R⁵ and a 5- or 6-membered ring containing atoms independently selected from the group consisting of C, N, O and S;

R⁵ and R⁶ are independently selected from, H, C₁₋₆alkyl, C₃₋₇cycloalkyl and aryl;

m is selected from 0, 1, 2, 3 or 4;

n is selected from 0, 1, 2, 3 or 4;

p is selected from 0, 1, 2, 3 or 4; and

a salt or hydrate thereof,

with the proviso that the compound is not:

4,4'-(1,2-piperazinediyl)di-antipyrine;

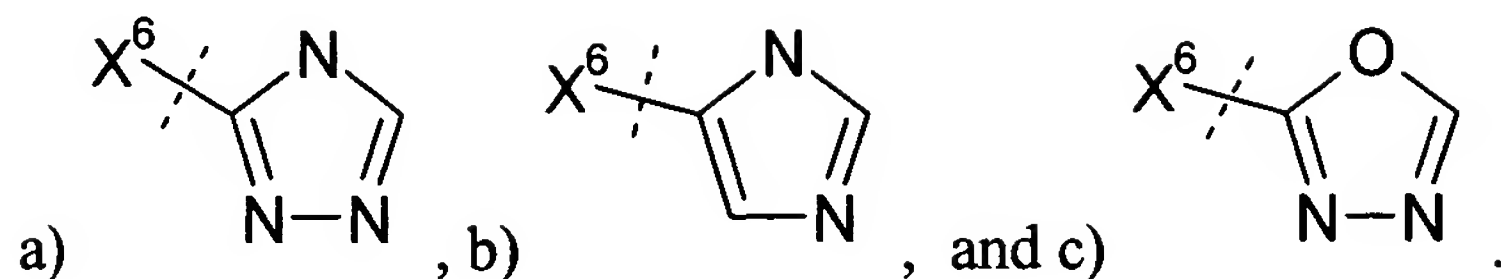
4,4'-(1,2-piperazinediyl)di-antipyrine dihydrochloride; or

4,4'-(1,2-piperazinediyl)di-antipyrine dipicrate;

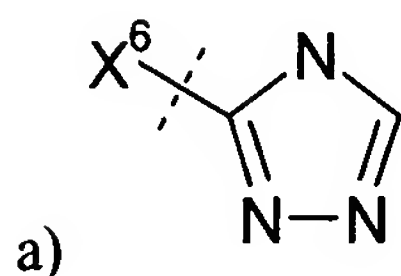
2. (Original) A compound according to claim 1 wherein m is selected from 1, 2, 3 or 4
3. (Original) A compound according to claim 1 wherein X^7 is C.
4. (Original) A compound according to claim 1 wherein X^5 is selected from the group consisting of $CR^4R^{4'}$, NR^4 , O, S, SO and SO_2 .
5. (Original) A compound according to claim 1 wherein X^3 is selected from the group consisting of N and O.
6. (Original) A compound according to claim 1 wherein P is aryl.
7. (Original) A compound according to claim 6 wherein P is phenyl.
8. (Original) A compound according to claim 7 wherein m is selected from the group consisting of 1 and 2.

9. (Original) A compound according to claim 1 wherein R^1 is selected from the group consisting of: halo, C_{1-6} alkylhalo, OC_{1-6} alkylhalo, C_{1-6} alkyl, OC_{1-6} alkyl, C_{1-6} alkylOR⁵, C_{0-6} alkylcyano, C_{0-6} alkylNR⁵R⁶.
10. (Original) A compound according to claim 9 wherein R^1 is selected from the group consisting of: Cl, F, Me, OMe, CF₃, OCF₃, and CN.
11. (Original) A compound according to claim 1 wherein X^2 is C.
12. (Original) A compound according to claim 11 wherein X^1 is N or CR⁴.
13. (Original) A compound according to claim 12 wherein when X^3 is O, X^4 is N and when X^3 is N, X^4 is O.
14. (Original) A compound according to claim 1 wherein X^2 is N.
15. (Original) A compound according to claim 14 wherein X^1 is N.
16. (Original) A compound according to claim 15 wherein X^3 is N and X^4 is N or CR⁴.

17. (Original) A compound according to claim 1 wherein X^6 is N.
18. (Original) A compound according to claim 12 wherein X^5 is selected from the group consisting of a bond, $CR^4R^{4'}$, NR^4 and O.
19. (Original) A compound according to claim 13 wherein X^5 is selected from the group consisting of a bond, O and NR^4 .
20. (Original) A compound according to claim 16 wherein X^5 is selected from the group consisting of O and CR^4 .
21. (Original) A compound according to claim 1 wherein R^4 is selected from the group consisting of: hydrogen, C_{1-6} alkyl, C_{1-6} alkylhalo and halo.
22. (Original) A compound according to claim 1 wherein Q is heteroaryl.
23. (Original) A compound according to claim 1 wherein Q is selected from the group consisting of:



24. (Original) A compound according to claim 23 wherein Q is



25. (Original) A compound according to claim 1 wherein R^2 and R^3 are independently selected from the group consisting of: C_{1-4} alkylhalo, C_{1-6} alkyl, C_{3-6} cycloalkyl, C_{0-6} alkylaryl and C_{0-6} alkylheteroaryl.

26. (Original) A compound according to claim 1 wherein A is selected from the group consisting of: hydrogen, hydroxyl, halo, C_{0-6} alkylcyano, C_{1-6} alkyl, $-OC_{1-6}$ alkyl, C_{1-6} alkylhalo, OC_{1-6} alkylhalo.

27. (Original) A compound according to claim 1 selected from:

4-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-piperidin-1-yl}-4-methyl-4H [1,2,4]triazol-3-yl)-pyridine

3-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-4-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-
morpholine

3-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-4-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-
[1,2,4]triazol-3-yl]-morpholine

3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-
yl)-morpholine

3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-
[1,2,4]triazol-3-yl]-morpholine

3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-
yl)-piperazine-1-carboxylic acid tert-butyl ester

2-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-1-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-
piperazine

2-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-methyl-1-(4-methyl-5-pyridin-4-yl-4H-
[1,2,4]triazol-3-yl)-piperazine

3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-
[1,2,4]triazol-3-yl]-piperazine-1-carboxylic acid tert-butyl ester

2-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-1-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-
[1,2,4]triazol-3-yl]-piperazine

2-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-1-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-
[1,2,4]triazol-3-yl]-4-methyl-piperazine

2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]-1-{5-[4-(difluoromethoxy)phenyl]-4-methyl-4H-1,2,4-triazol-3-yl}piperidine

4-(5-{2-[2-(3-chlorophenyl)-2H-tetrazol-5-yl]piperidin-1-yl}-4-methyl-4H-1,2,4-triazol-3-yl)pyridine

2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]-1-[5-(4-methoxyphenyl)-4-methyl-4H-1,2,4-triazol-3-yl]piperidine

[4-(5-{2-[2-(3-chlorophenyl)-2H-tetrazol-5-yl]piperidin-1-yl}-4-methyl-4H-1,2,4-triazol-3-yl)phenyl]dimethylamine

[4-(5-{2-[2-(3-Chloro-phenyl)-2H-tetrazol-5-yl]-piperidin-1-yl}-4-methyl-4H-[1,2,4]triazol-3-yl)-benzyl]-dimethyl-amine

{2-[4-(5-{2-[2-(3-Chloro-phenyl)-2H-tetrazol-5-yl]-piperidin-1-yl}-4-methyl-4H-[1,2,4]triazol-3-yl)-phenoxy]-ethyl}-dimethyl-amine

(R)-3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-morpholine

(S) 3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-morpholine

(R)-2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]-1-{5-[4-(difluoromethoxy)phenyl]-4-methyl-4H-1,2,4-triazol-3-yl}piperidine

(S)-2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]-1-{5-[4-(difluoromethoxy)phenyl]-4-methyl-4H-1,2,4-triazol-3-yl}piperidine

(R)-4-(5-{2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]piperidin-1-yl}-4-methyl-4H-1,2,4-triazol-3-yl)pyridine

(S)-4-(5-{2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]piperidin-1-yl}-4-methyl-4H-1,2,4-triazol-3-yl)pyridine

4-[5-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-pyrrolidin-1-yl}-4-cyclopropyl-4H-[1,2,4]triazol-3-yl)-pyridin-2-yl]-morpholine,

4-[5-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-pyrrolidin-1-yl}-4-methyl-4H-[1,2,4]triazol-3-yl)-pyridin-2-yl]-morpholine,

3-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-pyrrolidin-1-yl}-4-methyl-4H-[1,2,4]triazol-3-yl)-pyridine,

4-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-pyrrolidin-1-yl}-4-cyclopropyl-4H-[1,2,4]triazol-3-yl)-pyridine,

3-[5-(3-Chloro-phenyl)-[1,2,4]oxadiazol-3-yl]-4-(5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-morpholine,

3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-(4-cyclopropyl-5-pyridin-3-yl-4H-1,2,4-triazol-3-yl)morpholine,

3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-(4-cyclopropyl-5-pyridin-4-yl-4H-1,2,4-triazol-3-yl)morpholine,

3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-(4-methyl-5-pyridin-3-yl-4H-1,2,4-triazol-3-yl)morpholine,

3-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-4-[5-(6-methoxy-pyridin-3-yl)-4-methyl-4H-
[1,2,4]triazol-3-yl]-morpholine,

3-[3-(3-chlorophenyl)-1,2,4-oxadiazol-5-yl]-4-[5-(2-methoxypyridin-4-yl)-4-methyl-4H-1,2,4-
triazol-3-yl]morpholine,

3-[3-(3-chlorophenyl)-1,2,4-oxadiazol-5-yl]-4-[5-(2-methylpyridin-4-yl)-4-methyl-4H-1,2,4-
triazol-3-yl]morpholine,

3-[3-(3-chlorophenyl)-1,2,4-oxadiazol-5-yl]-4-[5-(5-fluoropyridin-3-yl)-4-methyl-4H-1,2,4-
triazol-3-yl]morpholine,

3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-[5-(5-fluoropyridin-3-yl)-4-methyl-4H-1,2,4-triazol-3-
yl]morpholine,

3-[3-(3-chlorophenyl)-1,2,4-oxadiazol-5-yl]-4-(4-methyl-5-pyridin-2-yl-4H-1,2,4-triazol-3-
yl)morpholine,

4-[5-(5-fluoropyridin-3-yl)-4-methyl-4H-1,2,4-triazol-3-yl]-3-[3-(3-iodophenyl)-1,2,4-oxadiazol-
5-yl]morpholine,

3-[3-(3-iodophenyl)-1,2,4-oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-1,2,4-triazol-3-
yl)morpholine,

3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-[5-(2-methylpyridin-4-yl)-4-methyl-4H-1,2,4-triazol-3-
yl]morpholine,

3-[2-(3-chlorophenyl)-2H-tetrazol-5-yl]-4-(4-methyl-5-pyridin-3-yl-4H-1,2,4-triazol-3-
yl)morpholine,

3-[2-(3-chlorophenyl)-2H-tetrazol-5-yl]-4-[5-(3,5-difluorophenyl)-4-methyl-4H-1,2,4-triazol-3-yl]morpholine,

3-(5-{2-[5-(3-chlorophenyl)isoxazol-3-yl]pyrrolidin-1-yl}-4-cyclopropyl-4H-1,2,4-triazol-3-yl)pyridine, and

4-(5-{2-[5-(3-chlorophenyl)isoxazol-3-yl]pyrrolidin-1-yl}-4-methyl-4H-1,2,4-triazol-3-yl)pyridine.

28. (Original) A pharmaceutical composition comprising as active ingredient a therapeutically effective amount of the compound according to any one of claims 1 to 26, in association with one or more pharmaceutically acceptable diluent, excipients and/or inert carrier.

29. (CANCELLED)

30. (Currently Amended) The compound according to ~~any one of claims 1 to 27~~ claim 1, for use in therapy.

31. (Currently Amended) The compound according to ~~any one of claims 1 to 27~~ claim 1, for use in treatment of mGluR 5 mediated disorders.

32. (Currently Amended) Use of the compound according to ~~any one of claims 1 to 27~~ claim 1, in the manufacture of a medicament for the treatment of mGluR 5 mediated disorders.

33. (Currently Amended) A method of treatment of mGluR 5 mediated disorders, comprising administering to a mammal, including man in need of such treatment, a therapeutically effective amount of the compound according to ~~any one of claims 1 to 27~~ claim 1.

34. (Original) The method according to claim 33, for use in treatment of neurological disorders.

35. (Original) The method according to claim 33, for use in treatment of psychiatric disorders.

36. (Original) The method according to claim 33, for use in treatment of chronic and acute pain disorders.

37. (Original) The method according to claim 33, for use in treatment of gastrointestinal disorders.

38. (Original) A method for inhibiting activation of mGluR 5 receptors, comprising treating a cell containing said receptor with an effective amount of the compound according to claim 1.